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# 1    **Assessing species vulnerability to climate change**

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## 60    **Abstract**

61            The effects of climate change on biodiversity are increasingly well documented, and many  
62    methods have been developed to assess species' vulnerability to climatic changes, both ongoing and  
63    projected in the coming decades. To minimize global biodiversity losses, conservationists need to  
64    identify those species that are likely to be most vulnerable to the impacts of climate change. In this  
65    review, we summarise different currencies used for assessing species' climate change vulnerability.  
66    We describe three main approaches used to derive these currencies (correlative, mechanistic and  
67    trait-based), and their associated data requirements, spatial and temporal scales of application and  
68    modelling methods. We identify strengths and weaknesses of the approaches and highlight the  
69    sources of uncertainty inherent in each approach that limit projection reliability. Finally, we provide  
70    guidance for conservation practitioners in selecting the most appropriate approach(es) for their  
71    planning needs and highlight priority areas for further assessments.

72

73           The Earth has warmed by about 0.74 °C in the last 100 years, and global mean temperatures  
74 are projected to increase further by 4.3 +/- 0.7 °C by 2100<sup>1</sup>. Agricultural expansion,  
75 overexploitation and invasive alien species introductions have been the main drivers of biodiversity  
76 loss in the recent past, but several lines of research suggest that climate change could become a  
77 prominent, if not leading cause of extinction over the coming century<sup>2</sup>, both via direct impacts on  
78 species and through synergies with other extinction drivers<sup>1,3</sup>. Species have already responded to  
79 recent climatic shifts<sup>4-8</sup>, and various attempts have been made to assess the potential risks to  
80 biodiversity posed by climate change over coming decades<sup>9-11</sup>.

81           To assess the threats to a species posed by climate change one must have information  
82 regarding its vulnerability, which is defined by the IPCC as ‘the predisposition to be adversely  
83 affected’<sup>12</sup>. Although there is currently no broad consensus in the scientific literature regarding the  
84 definition of ‘species’ vulnerability’, it is generally accepted that this is a function of both intrinsic  
85 and extrinsic factors<sup>13</sup>, and assessments often consider exposure, sensitivity and adaptability in  
86 combination<sup>13,14</sup>. Exposure is the magnitude of climatic variation in the areas occupied by the  
87 species<sup>15</sup>. Sensitivity, which is determined by traits that are intrinsic to species, is the ability to  
88 tolerate climatic variations, while adaptability is the inherent capacity of species to adjust to those  
89 changes<sup>14,15</sup>. Attempts at projecting the effects of climate change on species have used both  
90 different currencies (i.e. the range of measures used to assess species’ climate change vulnerability)  
91 and divergent approaches for identifying the most vulnerable taxa. Because of this lack of  
92 consensus by the conservation community, a formal comparative evaluation is necessary to guide  
93 sensible choices of the most appropriate technique(s) for assessing species’ vulnerability.

94           Here we provide the first comprehensive review of currencies and approaches that have been  
95 used to assess species’ vulnerability to climate change, based on a total of 97 studies published  
96 between 1996 and 2014 (with >70% of the studies published during the last five years). We  
97 describe the four dominant currencies of species’ climate change vulnerability assessments and

98 provide examples of how these have been applied. Three broad categories of approaches plus three  
99 combinations thereof were identified, and we describe each examining how they address  
100 uncertainties, and discuss their key limitations. Finally, we provide guidance for practitioners. Via  
101 these analyses, we aim to help conservationists select appropriate approaches for assessing species'  
102 vulnerability, such that climate change adaptation responses are as solidly based as possible.

103

#### 104 **Taxonomic and regional application of climate change vulnerability assessments of species**

105 We conducted a systematic literature search using ISI Web of Knowledge. Key-words were  
106 selected to identify studies on climate change (climate change\*, global warming\*, sea-level rise\*,  
107 elevated CO<sub>2</sub>\*, drought\*, cyclones\*, CO<sub>2</sub> concentration\*) impacts (population reduction\*, range  
108 changes\*, range shift\*, turnover\*, extinction risk\*, extinction probability\*) that led to vulnerability  
109 assessments (vulnerability\*, sensitivity\*, adaptability\*, exposure\*) based on different types of  
110 approaches (mechanistic\*, SDM\*, correlative\*, trait-based\*, criteria\*, niche models\*). We then  
111 selected the most representative papers (in terms of both spatial and temporal scales, and taxa).  
112 Studies differed widely in taxonomic coverage, birds being the most frequently considered taxon,  
113 followed by mammals and plants, while non-insect invertebrates being seldom assessed (Fig. 1).  
114 Additionally, spatial scales of application and authors' interpretations of the concept of  
115 vulnerability varied extensively. More than 60% of the studies were developed at local scale, while  
116 only 4% of the papers assessed species' vulnerability globally (Fig. 1). As a result, numerous  
117 species have been assessed in only part of their range and their estimates of vulnerability may  
118 therefore be unrealistic.

119 Many published studies have shown that life-history traits are more important than  
120 taxonomy and distribution in determining species vulnerability to climate change<sup>14</sup>. Traits that  
121 commonly make a species vulnerable to climate change include limited dispersal abilities<sup>14,16–18</sup>,

122 slow reproductive rates<sup>11,19</sup>, specialised habitat and dietary requirements<sup>14,20,21</sup>, restricted  
123 distribution and rarity<sup>14,22</sup>, and narrow physiological tolerances<sup>23–25</sup>, while potentially vulnerable  
124 habitats include intertidal areas, montane habitats, savannahs and grasslands<sup>25</sup>. Knowing what  
125 makes a species vulnerable and where vulnerable species are located can be very useful when  
126 practitioners need to assess the vulnerability of species for which only basic knowledge of their  
127 biology and ecology is available.

128         Studies conducted at a broad scale (regional, continental and global, almost 70% of the  
129 total), where used to derive a map of the areas with the greatest concentration of vulnerable species,  
130 according to an ecoregional classification (Fig. 2). For marine areas we performed a qualitative  
131 assessment (high, medium and low vulnerability, mostly based on Foden *et al.*<sup>14</sup>) because only a  
132 few marine taxa have been evaluated at broad scales and more than 80% of the species assessed are  
133 corals, while for terrestrial areas we were able to identify hotspots of vulnerable species as areas  
134 with high concentrations of vulnerable species (> 100), belonging to different taxonomic classes.  
135 These vulnerable areas, the Caribbean, the Amazon basin, Mesoamerica, eastern Europe through  
136 central and eastern Asia, the Mediterranean basin, the Himalayas, South-East Asia, North Africa,  
137 the Congo basin, tropical West Africa and Madagascar, should be a first priority for monitoring.  
138 However, over 70% of the studies we reviewed involved only three continents/subcontinents, with  
139 almost 33% of the studies in North America, 24% in Europe, and 14% in Australia (Fig. 3). By  
140 contrast, there is a paucity of studies in the most biodiverse tropical and subtropical regions of the  
141 world. Since climate change will act in concert with other threats, and habitat loss is predicted to  
142 severely affect biodiversity in developing countries<sup>26</sup>, it is essential to conduct studies in these data  
143 deficient areas.

144



## 145    **Currencies used to assess vulnerability: ‘WHAT’**

146            There is no standard way to assess a species’ vulnerability to climate change, and the type of  
147 information (e.g. range extent, population size) needed will determine which approaches are most  
148 appropriate.

149

## 150    **Distributional changes**

151            To assess climate change impacts on species, current and future distributions can be  
152 projected using either mechanistic or correlative niche models (both approaches are discussed  
153 below), which relate environmental conditions to species’ physiological responses or occurrence  
154 data, respectively. Several analyses have provided examples of species likely to suffer range  
155 reductions in the 21<sup>st</sup> century<sup>16,18</sup>. For example, Vieilleident *et al.*<sup>27</sup> predicted that the Malagasy  
156 baobab *Adansonia suarezensis* is likely to go extinct before 2080 due to an overall loss in suitable  
157 habitat. Changes in range size have usually been assessed by considering the climatic characteristics  
158 of current distributions and the projected distribution of these climatic conditions in future<sup>27,28</sup>.  
159 However, vulnerability might be exacerbated by other factors, including biotic interactions, reduced  
160 adaptive evolutionary response and dispersal ability. Several studies have incorporated dispersal  
161 ability into predictions of future range changes, either by contrasting scenarios of no dispersal with  
162 unlimited dispersal<sup>29–31</sup>, by estimating average or maximum potential dispersal distances<sup>16,18,24</sup>, or  
163 by explicitly simulating metapopulation dynamics including dispersal events<sup>32,33</sup>. For example,  
164 Schloss *et al.*<sup>18</sup> suggested that 87% of Western Hemisphere terrestrial mammals will likely  
165 experience a reduction in their climatically suitable area, with 20% of these species being  
166 particularly vulnerable due to their limited dispersal ability.

167

168

## 169    **Population changes**

170            A different set of modelling approaches uses predictions of population trends to inform risk  
171 assessments<sup>34</sup>. Quantified population changes can be based on direct observations, indices of  
172 abundance<sup>34–36</sup>, reporting rates used as proxies for abundance<sup>37</sup>, or they can be inferred from  
173 declines in extent of occupied or suitable habitat<sup>34,38</sup>. Examples of observed population declines  
174 within recent decades include long-distance avian migrants to Dutch forests, which have likely been  
175 driven principally by temperature changes in spring<sup>35</sup>. Also, a decrease in ice coverage has led to a  
176 reduction in polar bear (*Ursus maritimus*) numbers in the southern Beaufort Sea<sup>39</sup>. Some  
177 approaches to projecting future population sizes incorporate past population trends into mechanistic  
178 models<sup>39–41</sup>, and consider the effects of changes in model parameters (e.g. distribution patterns, life  
179 history, climatic conditions). This type of approach has also been applied to a population of  
180 American marten (*Martes americana*) in North America, where explicit population models have  
181 been used to simulate a 40% decline in the population due to climate change by 2055<sup>42</sup>.

182

## 183    **Extinction probability**

184            One synthesis estimated that between roughly 20 and 30% of species assessed are likely to  
185 be at increasingly high risk of extinction in the face of increasing global warming<sup>12</sup>. Extinction  
186 probability has been calculated for populations of species with known life-history characteristics,  
187 like the emperor penguin (*Aptenodytes forsteri*)<sup>41</sup>, Arizona cliffrose (*Purshia subintegra*)<sup>43</sup>, spring-  
188 summer chinook salmon (*Oncorhynchus tshawytscha*)<sup>44</sup> and polar bear (*Ursus maritimus*)<sup>39</sup>, by  
189 using Population Viability Analyses<sup>41,43</sup>, demographic models<sup>39,44,45</sup>, or evolutionary models<sup>46</sup>.  
190 These methodologies combine population fluctuations with changing environmental parameters in  
191 order to estimate extinction probability within a given time interval. For example, Fordham *et al.*<sup>45</sup>  
192 modelled the predicted abundance of the Iberian lynx (*Lynx pardinus*) under three climate scenarios

193 by integrating temperature and precipitation data, prey availability and management interventions,  
194 and predicted that climate change may drive this species to extinction within the next 50 years. This  
195 work relied upon a thorough understanding of the species' biology and of demographic dynamics  
196 related to extinction risk. However, as most species lack such detailed data, extinction risk due to  
197 climate change tends to be quantified only for better-known species.

198

## 199 **Vulnerability indices and other relative scoring systems**

200 Vulnerability indices are quantitative indicators of the relative vulnerability of species. The  
201 data derived from the currencies discussed above, and from trait-based vulnerability assessments  
202 (TVAs), can be used to obtain scores<sup>14</sup>, categories<sup>34</sup> or indices<sup>47</sup>, which are often easier for scientists  
203 and practitioners to interpret and use, in order to identify species at risk within their focal areas.  
204 Foden *et al.*<sup>14</sup>, for example, classified birds, amphibians and corals into two vulnerability categories  
205 (low or high). One limitation of indices and scores is that they do not provide any direct measures  
206 of the expected impact on species, i.e. they are not expressed in terms of any of the currencies  
207 otherwise used to assess species' vulnerability (e.g. range reductions, extinction probability,  
208 population decline).

209

## 210 **Approaches used to model species' vulnerability to climate change: 'HOW'**

211 Different approaches are used to assess species' vulnerability to climate change. These  
212 approaches can be placed in four classes: 1) correlative, 2) mechanistic, 3) trait-based, and 4)  
213 combined approaches.

214

215

## 216     **Correlative approaches**

217             Distributional changes are typically estimated through the use of correlative models that aim  
218     to represent the realized niche of a species<sup>48,49</sup>. Correlative models relate observed geographic  
219     distribution of a species to current climate; resultant models are then applied to climate projections  
220     to infer potential climatically-suitable areas for a given species in the future. Species' distribution  
221     can be presence-only data<sup>17,22</sup>, presence/absence<sup>50</sup> or abundance observations<sup>51</sup>, based either on  
222     fieldwork or specimen records<sup>22,52</sup>. Correlative models have been applied to species at scales  
223     ranging from local to global<sup>19,53</sup> (Fig. 1), and have been widely used to explore the vulnerability of  
224     vertebrates (including birds<sup>36,52,54</sup>, mammals<sup>17,28</sup>, amphibians<sup>30,50</sup>, fishes<sup>22,55</sup>), invertebrates<sup>14,56,57</sup>  
225     and plants<sup>27,58</sup>.

226             Correlative models have the advantage of being spatially explicit and they are applicable to  
227     a wide range of taxa at various spatial scales. However, there are a number of limitations and  
228     uncertainties associated with them (see Pearson *et al.*<sup>29</sup> and Wiens *et al.*<sup>59</sup> for detailed descriptions).  
229     Primary sources of uncertainty and potential errors can be divided into three broad classes: climatic,  
230     algorithmic, and biotic<sup>29,59</sup>. Climatic uncertainties, that apply to all types of approaches, may arise  
231     from general circulation models, which use different parameters and model structures to simulate  
232     future climate systems, and may produce different results irrespective of the assumed greenhouse  
233     gas emissions<sup>59,60</sup>. Climate models project future climate conditions at a coarser scale of resolution  
234     than that of data (biological and environmental) used to calibrate the correlative models<sup>49,59</sup>, and  
235     their outputs are thus often not sufficiently fine-scaled for modelling rare species or species with  
236     small geographic distributions<sup>49,50</sup>. Algorithmic uncertainties can arise from the differences in  
237     methods and models used to predict species' distribution (e.g. Generalized Additive Models,  
238     Maximum Entropy models), and from the selection of model predictors (e.g. mean annual  
239     temperature, annual precipitation; see<sup>61</sup>), which have shown great variability in both results and  
240     model performance. This range of uncertainties has been addressed by some by applying a variety

241 of different statistical methods and model structures, summarising predictions across all models to  
242 generate ensemble forecasts, e.g. model-averaged probability of presence and confidence intervals  
243 (see examples<sup>16,30,62</sup>). Biotic uncertainties may arise if the assumptions made about a species'  
244 biology are inappropriate. First, species' distributions are assumed to be in equilibrium with  
245 surrounding climates and these relationships are assumed to persist in the future<sup>56</sup>. Second, it is  
246 unknown how much of a species' fundamental niche, exclusively determined by the species'  
247 requirements and/or tolerances is represented by its currently realized niche, which is also  
248 determined by abiotic, biotic, geographic, historical and anthropogenic factors<sup>49</sup>. Moreover,  
249 correlative models for plants do not account for drivers such as changes in atmospheric CO<sub>2</sub>  
250 concentration, which influence plant growth and water use and can alter demographic processes  
251 sufficiently to drive ecosystem structural and functional changes<sup>63</sup>. Correlative models can also be  
252 used to predict future geographic distribution of a group of species in a given area and the results  
253 combined to create assessments of new community structures<sup>64</sup>. However, these models ignore  
254 community-assembly rules, as well as differences in the constraints and adaptability of individual  
255 species, and thus the resulting predicted species assemblages may be unrealistic<sup>62</sup>. Correlative  
256 models have been criticised by some authors because they lack mechanism and causality (e.g.  
257 see<sup>65</sup>), although there is increasing evidence that recent population trends have matched those  
258 expected from correlative model projections<sup>36</sup>.

259         The relatively large number of reliable occurrence points required to fit correlative models  
260 often precludes their use for assessments of poorly known species<sup>66</sup>. They are also less appropriate  
261 for species with cosmopolitan or limited geographic distributions (e.g. on small islands) since  
262 climate may not explain distributions or distributional changes. Despite these limitations, the  
263 majority of regional and global analyses to date are based on correlative approaches, since they can  
264 be relatively quick and cheap to apply<sup>67</sup> and occurrence data are available for a large number of  
265 taxa.

## 266    **Mechanistic approaches**

267            Mechanistic models require taxon-specific parameters that provide information on the  
268    behaviour of individuals and the mechanisms they develop to cope with changing climatic  
269    conditions. Mechanistic models are developed from laboratory and field observations of  
270    demographic rates, physiological tolerances<sup>41,68,69</sup>, competition and dispersal<sup>70</sup>, diseases and  
271    predation<sup>71</sup>, as well as from energy balance equations<sup>72</sup>. Measures of vulnerability derived from  
272    these models are typically expressed in terms of probability of extinction, whether of discrete  
273    populations or entire species. Mechanistic approaches often focus on a single species of  
274    conservation interest (e.g. rare or threatened species)<sup>39,41</sup>, since methods used to collect detailed data  
275    on species physiology, which are essential to parameterise such models, are costly and time-  
276    consuming. Some studies exist involving this type of modelling that do not involve a specific taxon  
277    but rather provide general theoretical frameworks to predict effects of climate change on plants<sup>10</sup>,  
278    terrestrial ectotherms<sup>68</sup> and generic species<sup>9,10</sup>, highlighting major determinants of extinction risk in  
279    a changing environment and providing recommendations for future research needs. Some  
280    mechanistic models (e.g. incidence function models, age-structured metapopulation models) may be  
281    used to explain metapopulation dynamics in the presence of climate change by estimating extinction  
282    and colonization rates as functions of habitat suitability<sup>73</sup>, prey availability or management  
283    actions<sup>45</sup>. Other mechanistic models consider the changes in vegetation distribution and dynamics  
284    using both bioclimatic and physiological parameters of groups of species (e.g. plant functional  
285    types)<sup>74</sup>.

286            Mechanistic niche models utilise species' functional traits, physiological tolerances and  
287    energy and mass exchanges to represent the fundamental niche of a species<sup>75</sup>. Key functional traits  
288    (e.g. morphology, physiology, behaviour) and spatial habitat data (e.g. climate, vegetation cover,  
289    topography, bathymetry) are used to assess individual fitness<sup>75,76</sup>. Such models are considered by  
290    some authors to be more robust and theoretically defensible than correlative models for predicting

species' responses to climate change<sup>75</sup>. Compared to the realized niche modelled via correlative approaches, the mechanistically modelled fundamental niche provides a better approximation of the climatic space in which an organism can exist, including areas that have, or may, become newly suitable<sup>75,76</sup>. In addition, these models permit explicit consideration of important biological factors like evolutionary changes and physiological responses.

Extensive application of mechanistic niche models is precluded by the fact that they require detailed data that are lacking for most species. The main sources of uncertainty in mechanistic models relate to model parameters (e.g. population abundance, which may be underestimated depending on the method used to collect the data and the ability of the observer to detect the species), and to combining data collected at different spatial resolutions<sup>23</sup>. Moreover, these models usually do not account for non-climatic threats to dispersal or for biotic interactions<sup>48</sup>.

### **Trait-based vulnerability assessment approaches**

TVAs use species' biological characteristics as predictors of extinction risk due to climate change<sup>13,14</sup>, often in combination with estimates of exposure. Methods typically involve selecting traits related to sensitivity (e.g. typically describing ecological specialization, inter-specific interactions) and adaptability (i.e. dispersal and phenotypic adaptability<sup>14,77,78</sup>) and scoring each according to observations or expert judgment<sup>79,80</sup>. For example, Gardali *et al.*<sup>78</sup> quantified the vulnerability of Californian birds by scoring sensitivity and exposure for each taxon. They used information from published literature to assign a sensitivity score to four intrinsic species' characteristics (dispersal ability, migratory status, habitat specialization and physiological tolerances), and then combined sensitivity and exposure scores to generate a climate vulnerability index.

314 TVAs are being used increasingly by conservation organizations and management agencies  
315 because they permit a relatively rapid assessment for multiple species, which can be used to  
316 prioritize conservation planning and implementation of adaptation schemes. Moreover, TVAs are  
317 sometimes considered easier to use by practitioners because they do not require extensive  
318 knowledge of modelling techniques, even if their applicability is limited to a specific area and to  
319 cases where relevant data on species' traits are available (see<sup>81</sup>).

320 Drawbacks with TVAs are that precise vulnerability thresholds associated with each trait are  
321 often unknown, necessitating selection of arbitrary, relative thresholds for categories of higher or  
322 lower extinction risk. Traits are often weighted equally<sup>20</sup> even though some characteristics are likely  
323 to be more important than others in determining climate change vulnerability. Subject to the  
324 challenges of score-based systems, it is not possible to compare vulnerability between taxonomic  
325 groups for which different sets of traits may have been used in the TVA. Moreover, different TVAs  
326 applied to the same species do not always yield congruent results<sup>82</sup>. The most common sources of  
327 uncertainty in TVAs stem from the choice of traits included in assessments, parameterisation of  
328 thresholds of associated vulnerability, and from gaps in knowledge of individual species'  
329 characteristics<sup>14,83</sup>. For example, dispersal distance is one of TVA's most important and  
330 conservation-informative traits, yet estimates are currently available for few animal species. Some  
331 studies have attempted to provide dispersal estimates<sup>16,18,84</sup>, but inevitable uncertainties arise from  
332 models and parameters. Uncertainty is usually incorporated as a confidence score based on expert  
333 opinion. Such score can be provided for each trait<sup>78</sup>, for each stage of the assessment<sup>83</sup>, or for the  
334 overall assessment<sup>78</sup>. Alternatively, some authors rank missing trait data under best- or worst-case  
335 scenarios<sup>14,80</sup>, by assuming optimistic and pessimistic extreme values.

336



## 337    **Combined approaches**

338            There is a growing consensus on the benefits of using approaches that combine different  
339 types of models and data<sup>32,40</sup>. Here we discuss the three most common combined approaches,  
340 criteria-based, mechanistic-correlative and correlative-TVA.

341

## 342    **Criteria-based approaches**

343            Criteria-based approaches have been used to combine observed or projected demographic  
344 trends (e.g. population increases or decreases) with intrinsic characteristics of species (e.g.  
345 generation length), to classify species into threat categories based on the risks posed by climate  
346 change. Climate-attributed changes in species' geographic ranges, often derived from correlative  
347 models, are assessed against quantitative thresholds<sup>34,38,83,85</sup>. These assessments often use the IUCN  
348 Red List categories and criteria ([www.iucnredlist.org](http://www.iucnredlist.org))<sup>38,85</sup> or draw inspiration from them<sup>83</sup>.

349            One advantage of criteria-based approaches is that they can be applied to large numbers of  
350 species worldwide<sup>86</sup>. They are important for assessing the conservation status of species threatened  
351 by climate change since they simultaneously account for several factors known to affect the relative  
352 extinction risk (e.g. declines in the extent of occurrence, reduction in population size). Furthermore,  
353 by using quantitative thresholds to predict relative extinction risk, it is possible to make  
354 comparisons between past, current and future conservation status of species. Approaches based on  
355 the IUCN Red List require a consistent adoption of thresholds and criteria<sup>87</sup>; however, these are  
356 sometimes arbitrarily modified (e.g. to temporal and spatial scales and spatial resolution), thereby  
357 reducing the comparability and interpretability of the results<sup>87</sup>. Pearson *et al.*<sup>88</sup> identified factors that  
358 predispose a selection of North American herpetiles to high extinction risk due to climate change,  
359 and concluded that most important factors are already incorporated into extinction risk assessments  
360 for the IUCN Red List.

361     **Mechanistic-correlative and mechanistic-correlative-TVA approaches.**

362             In mechanistic-correlative approaches, outputs of correlative models are incorporated into  
363     demographic models to calculate spatial structure of populations<sup>45</sup>, whose dynamics are then  
364     modelled mechanistically. This combination is useful, for example, in predicting how distribution  
365     patterns influence the viability of populations under a changing climate<sup>32,40</sup>. Furthermore, some  
366     studies have integrated life-history characteristics into models to produce more accurate projections  
367     of species' responses to climate change. Keith *et al.*<sup>32</sup> assessed extinction risk for plant species in  
368     South African fynbos under stable and changing climatic conditions. The authors linked the outputs  
369     of correlative models with a demographic metapopulation model, and considered their interactions  
370     with fire tolerances and dispersal abilities. In this way, they dealt with both habitat changes and  
371     population dynamics simultaneously in their assessments.

372

373     **Correlative-TVA approaches**

374             Other combined approaches integrate species characteristics and species distribution models  
375     by incorporating species traits to refine distribution projections made using correlative  
376     models<sup>16,18,31,89</sup>, or by integrating correlative model outputs into trait-based assessments<sup>21,83</sup>. In the  
377     first approach, traits like dispersal ability and generation length have been usefully applied to refine  
378     range dynamics<sup>16,90</sup>. For example, Barbet-Massin *et al.*<sup>16</sup> used natal dispersal and generation length  
379     to predict the breeding distribution of European birds under climate and land-use changes. The  
380     authors predicted a 10% reduction of future species richness assuming unlimited dispersal and a  
381     25% reduction by using natal dispersal.

382             In the second type of approach, the outputs of correlative models are used to estimate  
383     exposure to climate change and identify areas, which might become suitable in the future, even if  
384     they fall outside a species' current range. By linking exposure, estimated with correlative models,

385 with sensitivity and adaptability assessed with TVAs, a vulnerability index can be calculated that  
386 accounts for both intrinsic and extrinsic factors (e.g.<sup>83</sup>).

387

## 388 **Guidance for selecting climate change vulnerability assessment approaches**

389 Ideally, practitioners should assess the vulnerability of populations or species to climate  
390 change using a variety of methods, with greatest predictive confidence conferred where models are  
391 in agreement. The choice of the approach is entirely dependent on conservation goals, which are  
392 often vague and not clearly defined, and on the data available (Box 1). Relying on these broad  
393 goals, practitioners need to identify definable and measurable objectives<sup>91</sup>, in terms of temporal,  
394 spatial and taxonomic scales. In Table 1 we identify different examples of objectives against each  
395 approach and below provide two exemplary goals and identify the associated methodologies to  
396 reach them.

397

## 398 **Estimating extinction risk**

399 When deriving estimates of extinction risk of species is the goal, both mechanistic and  
400 correlative models can provide appropriate results. The most effective way to predict extinction risk  
401 of species under climate change is to combine demographic data (e.g. population trends, survival,  
402 fertility) with changing environmental factors (e.g. precipitation, sea ice extent), and then project  
403 these changes into the future<sup>41,43</sup>. For example, Jenouvrier *et al.*<sup>41</sup> used a mechanistic model, which  
404 combined demographic and climatic data, to project a > 35% probability of extinction for the  
405 emperor penguins (*Aptenodytes forsteri*) in Antarctica by 2100 in response to projected sea ice  
406 changes.

407 Another way of inferring the extinction risk of species is to use a decline in suitable area as a  
408 proxy for population decline<sup>38,92,93</sup>, providing the relationship between the two can be assumed to  
409 remain constant. Correlative models can be used to project range changes into the future; this would

410 allow classifying the species into one of IUCN Red List categories. Levinsky *et al.*<sup>93</sup>, for example,  
411 demonstrated that the proportion of European mammals that are forecast to become extinct by 2100  
412 can vary from 1 to 9%, depending on the magnitude of predicted climatic changes and the ability of  
413 species to migrate.

414

#### 415 **Prioritization of actions**

416 Climate change adaptation strategies require creating a link between an explicitly stated  
417 expectation about the way global warming could affect species, habitats, or even people, to clear  
418 objectives and actions that would best address those climate impacts<sup>94</sup>. Conservation decision-  
419 making is about prioritizing actions to satisfy conservation objectives for a set of species and  
420 areas<sup>95</sup>. It is not possible to make conservation interventions for all species, and prioritization  
421 exercises are needed to determine which actions to focus on to protect species<sup>96</sup>. Given the high  
422 levels of uncertainty and complexity in modelling impacts, we highlight that reprioritizing or even  
423 abandoning actions which benefit some species over others should be done with great caution.

424 Where site-scale conservation is the focus (e.g. in a protected area), correlative models are  
425 able to identify species for which the area may be suitable in the future, thereby allowing managers  
426 to prepare for potentially novel species assemblages and plan appropriate conservation actions (e.g.  
427 predator and invasive species control). For example, Hole *et al.*<sup>54</sup> used correlative models to assess  
428 species turnover in a network of Important Bird Areas in Africa, and provided generic guidance on  
429 the types of conservation actions (e.g. translocation, habitat restoration, disturbance-regime  
430 management) that might be most appropriate for individual sites.

431 For a regional-scale focus, identifying the bioclimatic space where species could persist and  
432 the areas of relatively unchanged climate within this space may facilitate species persistence during  
433 periods of climatic stress. Spatially explicit projections from correlative and mechanistic niche  
434 models allow identification of these sites. For example, Maschinski *et al.*<sup>43</sup> used a mechanistic

435 approach to identify potential climatic refugia for an endemic plant species (*Purshia subintegra*) of  
436 Arizona. This study showed that in situ manipulation and introductions at northern latitudes are  
437 priority actions necessary to prevent the extinction of this rare and endangered species.

438         Where the focus is on particular species, trait-based and mechanistic approaches are likely to  
439 deliver insights into the specific mechanism(s) of impact (e.g. increased competition, loss of  
440 mutualisms, disruption of cues, disease)<sup>14</sup>, allowing targeted interventions both to decrease species'  
441 sensitivity (e.g. disease treatment, predator control) and to increase their adaptive capacity (e.g.  
442 genetic management, improved landscape permeability, translocation)<sup>75</sup>. Indices calculated with  
443 trait-based approaches can facilitate grouping taxa by their relative risk to climatic changes, which  
444 help identify adaptation strategies that could benefit multiple species<sup>77</sup>. For example, Moyle *et. al*<sup>80</sup>,  
445 who assessed Californian freshwater fishes according to their life-history characteristics, classified  
446 species that were heavily dependent on human intervention as highly vulnerable to climate change,  
447 and highlighted the need for conservation actions such as management of barriers, special flows and  
448 removal of alien species to allow population persistence.

449

## 450 **Conclusions**

451         This review of climate change vulnerability assessment approaches suggests that, in general,  
452 a correlative approach is appropriate when the only data available are those on species' occurrence,  
453 in particular for reconstructing the paleoclimatic niche of fossil species or projecting their future  
454 climatic suitable area, from local to global scales. On the other hand, mechanistic models have the  
455 greatest power to assess extinction probability driven by climate change, identify conservation  
456 actions and evaluate the potential effectiveness of management interventions, but they are limited to  
457 few terrestrial species. Therefore, they are usually employed when the focus is on a well-studied  
458 species of particular conservation interest (e.g. species threatened, keystone, flagship or umbrella),  
459 for which detailed physiologic and/or demographic data are available. Trait-based approaches are

460 less resource-intensive and therefore more widely used. This method is ideal to help non-GIS  
461 experts develop regional assessments and to identify conservation priorities in the absence of  
462 specific data on species' distribution.

463 Validation of the accuracy and precision of vulnerability assessment approaches, through  
464 comparison of model projections with a globally coordinated observation effort, is essential for  
465 improving projections of the impacts of climate change on species. Use of paleoecological evidence  
466 of past species' responses to climatic variation in conjunction with matching paleoclimatic data can  
467 provide an opportunity to test the assessments<sup>97,98</sup>. Observations of recent responses to climate  
468 change are another useful tool to test reliability of model predictions against current observations.  
469 However, quantifying the ability of models to provide reliable range shift projections or population  
470 changes is still challenging, since they are often difficult to validate across time and space<sup>97</sup>. One  
471 key issue is the debate on modelling the realized vs. the fundamental niche<sup>48,49,79</sup>. Both the lack of  
472 equilibrium between species and climate, and the difficulty of isolating the effects of climatic  
473 changes on a species' range from those of other threats<sup>97</sup>, can lead to changes in the realized niche  
474 of a species (usually modelled mechanistically). On the other hand, correlative approaches attempt  
475 to model the fundamental niche of a species, but they use data from the realized niche<sup>48</sup>. This can  
476 lead to spurious correlations between species' occurrence and climate and thus hinder model  
477 validation as well as casting doubts on model accuracy<sup>48</sup>. For example, a species may not respond to  
478 climate only because other factors (e.g. competitive exclusion, predation) are confounding the  
479 response<sup>99</sup>. Additionally, when comparing past and current distribution to validate models or TVAs,  
480 a big challenge is to find accurate information on species' historic distribution and population  
481 trends. Addressing all of these issues should lead to better conservation decision-making.

482 A glaring oversight in almost all studies is that they only focused on the direct impacts of  
483 climate change. Indirect impacts within biological communities, as well as changes in human use of  
484 natural resources are going to have substantial, complex, and often multiplicative impacts on

485 species<sup>36,100</sup>. Thus, many current assessments are blind to the fact that the interactions between  
486 current threats and climate change are likely to be profound<sup>3</sup>. Moreover, the growing human  
487 population will itself be increasingly affected by climate change, with human adaptation responses  
488 likely to result in substantial and negative impacts on biodiversity<sup>100</sup>. Assessments of future impacts  
489 of climate change need to take these factors into account.

490

## 491    **References**

- 492    1.     Stocker, T. F., D. Qin, and G. K. Plattner. "Climate Change 2013: The Physical Science Basis."  
493        Working Group I Contribution to the Fifth Assessment Report of the Intergovernmental Panel on  
494        Climate Change. Summary for Policymakers (IPCC, 2013) (2013).
- 495    2.     Thomas, C.D. *et al.* Extinction risk from climate change. *Nature* **427**, 145–148 (2004).
- 496    3.     Mantyka-Pringle, C.S., Martin, T.G. & Rhodes, J.R. Interactions between climate and habitat loss  
497        effects on biodiversity: a systematic review and meta-analysis. *Glob. Chang. Biol.* **18**, 1239–1252  
498        (2012).
- 499    4.     Sinervo, B. *et al.* Erosion of lizard diversity by climate change and altered thermal niches. *Science*  
500        **328**, 894–9 (2010).
- 501    5.     Sheridan, J.A. & Bickford, D. Shrinking body size as an ecological response to climate change. *Nat.*  
502        *Clim. Change.* **1**, 401–406 (2011).
- 503    6.     Réale, D., McAdam, A.G., Boutin, S. & Berteaux, D. Genetic and plastic responses of a northern  
504        mammal to climate change. *Proc. Biol. Sci.* **270**, 591–6 (2003).
- 505    7.     Ockendon, N. *et al.* Mechanisms underpinning climatic impacts on natural populations: altered  
506        species interactions are more important than direct effects. *Glob. Change Biol.* (2014).  
507        doi:10.1111/gcb.12559
- 508    8.     Auer, S.K. & King, D.I. Ecological and life-history traits explain recent boundary shifts in elevation  
509        and latitude of western North American songbirds. *Glob. Ecol. Biogeogr.* (2014).  
510        doi:10.1111/geb.12174
- 511    9.     Chevin, L.-M., Lande, R. & Mace, G. M. Adaptation, plasticity, and extinction in a changing  
512        environment: towards a predictive theory. *PLoS Biol.* **8**, e1000357 (2010).
- 513    10.    Midgley, G.F. *et al.* BioMove - an integrated platform simulating the dynamic response of species to  
514        environmental change. *Ecography.* **33**, 612–616 (2010).
- 515    11.    Chessman, B. C. Identifying species at risk from climate change: Traits predict the drought  
516        vulnerability of freshwater fishes. *Biol. Conserv.* **160**, 40–49 (2013).
- 517    12.    Oppenheimer, M. *et al.* *Climate Change 2014: Impacts, Adaptation, and Vulnerability*. Contribution  
518        of working group II to the fifth assessment report of the intergovernmental panel on climate change,  
519        Ch. 19. 1–107 (2014).
- 520    13.    Williams, S.E., Shoo, L.P., Isaac, J.L., Hoffmann, A.A. & Langham, G. Towards an integrated  
521        framework for assessing the vulnerability of species to climate change. *PLoS Biol.* **6**, 2621–6 (2008).
- 522    14.    Foden, W.B. *et al.* Identifying the World's Most Climate Change Vulnerable Species: A Systematic  
523        Trait-Based Assessment of all Birds, Amphibians and Corals. *PLoS One* **8**, e65427 (2013).
- 524    15.    Dawson, T.P., Jackson, S.T., House, J.I., Prentice, I.C. & Mace, G.M. Beyond predictions:  
525        biodiversity conservation in a changing climate. *Science* **332**, 53–8 (2011).
- 526    16.    Barbet-Massin, M., Thuiller, W. & Jiguet, F. The fate of European breeding birds under climate,  
527        land-use and dispersal scenarios. *Glob. Change Biol.* **18**, 881–890 (2012).



- 528 17. Hughes, A.C., Satasook, C., Bates, P.J.J., Bumrungsri, S. & Jones, G. The projected effects of  
529 climatic and vegetation changes on the distribution and diversity of Southeast Asian bats. *Glob.*  
530 *Change Biol.* **18**, 1854–1865 (2012).
- 531 18. Schloss, C.A., Nuñez, T.A. & Lawler, J.J. Dispersal will limit ability of mammals to track climate  
532 change in the Western Hemisphere. *PNAS* **2012**, (2012).
- 533 19. Ameca y Juárez, E.I., Mace, G.M., Cowlishaw, G., Cornforth, W.A. & Pettoirelli, N. Assessing  
534 exposure to extreme climatic events for terrestrial mammals. *Conserv. Lett.* **6**, 145–153 (2013).
- 535 20. Laidre, K.L. *et al.* Quantifying the sensitivity of Arctic marine mammals to climate-induced habitat  
536 change. *Ecol. Appl.* **18**, S97–125 (2008).
- 537 21. Garnett, S. *et al.* *Climate change adaptation strategies for Australian birds*. 109 (2013).
- 538 22. Yu, D. *et al.* Global climate change will severely decrease potential distribution of the East Asian  
539 coldwater fish *Rhynchocypris oxycephalus* (Actinopterygii, Cyprinidae). *Hydrobiologia* **700**, 23–32  
540 (2013).
- 541 23. Buckley, L.B. The range implications of lizard traits in changing environments. *Glob. Ecol. Biogeogr.*  
542 452–464 (2010). doi:10.1111/j.1466-8238.2010.00538.x
- 543 24. Iverson, L.R., Prasad, A.M., Matthews, S.N. & Peters, M.P. Lessons Learned While Integrating  
544 Habitat, Dispersal, Disturbance, and Life-History Traits into Species Habitat Models Under Climate  
545 Change. *Ecosystems* **14**, 1005–1020 (2011).
- 546 25. Crossman, N.D., Bryan, B.A. & Summers, D.M. Identifying priority areas for reducing species  
547 vulnerability to climate change. *Divers. Distrib.* **18**, 60–72 (2012).
- 548 26. Visconti, P. *et al.* Future hotspots of terrestrial mammal loss. *Philos. Trans. R. Soc. London - Ser. B*  
549 *Biol. Sci.* **366**, 2693–2702 (2011).
- 550 27. Vieilledent, G., Cornu, C., Cuní Sanchez, A., Leong Pock-Tsy, J.-M. & Danthu, P. Vulnerability of  
551 baobab species to climate change and effectiveness of the protected area network in Madagascar:  
552 Towards new conservation priorities. *Biol. Conserv.* **166**, 11–22 (2013).
- 553 28. Songer, M., Delion, M., Biggs, A. & Huang, Q. Modeling Impacts of Climate Change on Giant Panda  
554 Habitat. *Int. J. Ecol.* **2012**, 1–12 (2012).
- 555 29. Pearson, R.G. *et al.* Model-based uncertainty in species range prediction. *J. Biogeogr.* **33**, 1704–1711  
556 (2006).
- 557 30. Carvalho, S.B., Brito, J.C., Crespo, E.G., Watts, M.E. & Possingham, H.P. Conservation planning  
558 under climate change: Toward accounting for uncertainty in predicted species distributions to  
559 increase confidence in conservation investments in space and time. *Biol. Conserv.* **144**, 2020–2030  
560 (2011).
- 561 31. Tuanmu, M.-N. *et al.* Climate-change impacts on understorey bamboo species and giant pandas in  
562 China's Qinling Mountains. *Nat. Clim. Change.* **3**, 249–253 (2012).
- 563 32. Keith, D.A. *et al.* Predicting extinction risks under climate change: coupling stochastic population  
564 models with dynamic bioclimatic habitat models. *Biol. Lett.* **4**, 560–3 (2008).

- 565 33. Fordham, D.A., Akçakaya, H.R., Araújo, M.B., Keith, D.A. & Brook, B.W. Tools for integrating  
566 range change, extinction risk and climate change information into conservation management.  
567 *Ecography* **36**,001-009 (2013).
- 568 34. UCN Standards and Petitions Subcommittee. Guidelines for Using the IUCN Red List Categories and  
569 Criteria. Version 10. Prepared by the Standards and Petitions Subcommittee (2013); available at  
570 <http://www.iucnredlist.org/documents/RedListGuidelines.pdf>.
- 571 35. Both, C. *et al.* Avian population consequences of climate change are most severe for long-distance  
572 migrants in seasonal habitats. *Proc. Biol. Sci.* **277**, 1259–66 (2010).
- 573 36. Gregory, R.D. *et al.* An indicator of the impact of climatic change on European bird populations.  
574 *PLoS One* **4**, e4678 (2009).
- 575 37. Huntley, B., Altwegg, R., Barnard, P., Collingham, Y.C. & Hole, D.G. Modelling relationships  
576 between species spatial abundance patterns and climate. *Glob. Ecol. Biogeogr.* **21**, 668–681 (2012).
- 577 38. Nenzén, H.K. & Araújo, M.B. Choice of threshold alters projections of species range shifts under  
578 climate change. *Ecol. Modell.* **222**, 3346–3354 (2011).
- 579 39. Hunter, C.M. *et al.* Climate change threatens polar bear populations: a stochastic demographic  
580 analysis. *Ecology* **91**, 2883–2897 (2010).
- 581 40. Anderson, B.J. *et al.* Dynamics of range margins for metapopulations under climate change. *Proc.*  
582 *Biol. Sci.* **276**, 1415–20 (2009).
- 583 41. Jenouvrier, S. *et al.* Demographic models and IPCC climate projections predict the decline of an  
584 emperor penguin population. *Proc. Natl. Acad. Sci.* **106**, 11425–11425 (2009).
- 585 42. Carroll, C. Interacting effects of climate change, landscape conversion, and harvest on carnivore  
586 populations at the range margin: marten and lynx in the northern Appalachians. *Conserv. Biol.* **21**,  
587 1092–104 (2007).
- 588 43. Maschinski, J., Baggs, J.E., Quintana-Ascencio, P.F. & Menges, E.S. Using Population Viability  
589 Analysis to Predict the Effects of Climate Change on the Extinction Risk of an Endangered  
590 Limestone Endemic Shrub, Arizona Cliffrose. *Conserv. Biol.* **20**, 218–228 (2006).
- 591 44. Thompson, L.C. *et al.* Water Management Adaptations to Prevent Loss of Spring-Run Chinook  
592 Salmon in California under Climate Change. *J. Water Resour. Plan. Manag.* **138**, 465–478 (2012).
- 593 45. Fordham, D.A. *et al.* Adapted conservation measures are required to save the Iberian lynx in a  
594 changing climate. *Nat. Clim. Change* **3**, 899–903 (2013).
- 595 46. Vedder, O., Bouwhuis, S. & Sheldon, B. C. Quantitative Assessment of the Importance of Phenotypic  
596 Plasticity in Adaptation to Climate Change in Wild Bird Populations. *PLoS Biol.* **11**, e1001605  
597 (2013).
- 598 47. Young, B. E., *et al.* Rapid assessment of plant and animal vulnerability to climate change. Pages 129-  
599 150 in *Wildlife Conservation in a Changing Climate*, (eds. Brodie, J., Post, E., & Doak, D. University  
600 of Chicago Press, Chicago, IL, 2012) (2012).
- 601 48. Pearson, R.G. & Dawson, T.P. Predicting the impacts of climate change on the distribution of  
602 species: are bioclimate envelope models useful? *Glob. Ecol. Biogeogr.* **12**, 361–371 (2003).

- 603 49. Guisan, A. & Thuiller, W. Predicting species distribution: offering more than simple habitat models.  
604 *Ecol. Lett.* **8**, 993–1009 (2005).
- 605 50. Lawler, J.J., Shafer, S.L., Bancroft, B.A. & Blaustein, A. R. Projected climate impacts for the  
606 amphibians of the Western hemisphere. *Conserv. Biol.* **24**, 38–50 (2009).
- 607 51. Howard, C., Stephens, P.A., Pearce-Higgins, J.W., Gregory, R.D. & Willis, S.G. Improving species  
608 distribution models: the value of data on abundance. *Methods Ecol. Evol.* (2014). doi:10.1111/2041-  
609 210X.12184
- 610 52. Garcia, R.A., Burgess, N.D., Cabeza, M., Rahbek, C. & Araújo, M.B. Exploring consensus in 21st  
611 century projections of climatically suitable areas for African vertebrates. *Glob. Change Biol.* **18**,  
612 1253–1269 (2012).
- 613 53. Watson, J.E.M., Iwamura, T. & Butt, N. Mapping vulnerability and conservation adaptation strategies  
614 under climate change. *Nat. Clim. Change* **3**, 989–994 (2013).
- 615 54. Hole, D.G. *et al.* Toward a management framework for networks of protected areas in the face of  
616 climate change. *Conserv. Biol.* **25**, 305–15 (2011).
- 617 55. Jeschke, J.M. & Strayer, D.L. Usefulness of bioclimatic models for studying climate change and  
618 invasive species. *Ann. N. Y. Acad. Sci.* **1134**, 1–24 (2008).
- 619 56. Harrison, P.A., Berry, P.M., Butt, N. & New, M. Modelling climate change impacts on species'  
620 distributions at the European scale: implications for conservation policy. *Environ. Sci. Policy* **9**, 116–  
621 128 (2006).
- 622 57. Sánchez-Fernández, D., Lobo, J.M. & Hernández-Manrique, O.L. Species distribution models that do  
623 not incorporate global data misrepresent potential distributions: a case study using Iberian diving  
624 beetles. *Divers. Distrib.* **17**, 163–171 (2011).
- 625 58. Cole, K.L. *et al.* Past and ongoing shifts in Joshua tree distribution support future modeled range  
626 contraction. *Ecol. Appl.* **21**, 137–49 (2011).
- 627 59. Wiens, J.A., Stralberg, D., Jongsomjit, D., Howell, C.A. & Snyder, M.A. Niches, models, and climate  
628 change: assessing the assumptions and uncertainties. *Proc. Natl. Acad. Sci. U. S. A.* **106 Suppl**,  
629 19729–36 (2009).
- 630 60. Bagchi, R. *et al.* Evaluating the effectiveness of conservation site networks under climate change:  
631 accounting for uncertainty. *Glob. Change Biol.* **19**, 1236–48 (2013).
- 632 61. Heikkinen, R.K. *et al.* Methods and uncertainties in bioclimatic envelope modelling under climate  
633 change. *Prog. Phys. Geogr.* **30**, 751–777 (2006).
- 634 62. Guisan, A. & Rahbek, C. SESAM - a new framework integrating macroecological and species  
635 distribution models for predicting spatio-temporal patterns of species assemblages. *J. Biogeogr.* **38**,  
636 1433–1444 (2011).
- 637 63. Higgins, S.I. & Scheiter, S. Atmospheric CO<sub>2</sub> forces abrupt vegetation shifts locally, but not globally.  
638 *Nature* **488**, 209–12 (2012).
- 639 64. Dubuis, A. *et al.* Predicting spatial patterns of plant species richness: a comparison of direct  
640 macroecological and species stacking modelling approaches. *Divers. Distrib.* **17**, 1122–1131 (2011).

- 641 65. Tyre, A. J., Possingham, H. P., & Lindenmayer, D. B. Inferring process from pattern: can territory  
642 occupancy provide information about life history parameters?. *Ecological Applications*, **11**, 1722-  
643 1737 (2001).
- 644 66. Boitani, L. *et al.* What spatial data do we need to develop global mammal conservation strategies?  
645 *Philos. Trans. R. Soc. Lond. B. Biol. Sci.* **366**, 2623–32 (2011).
- 646 67. Peterson, A.T. & Martínez-Meyer, E. Geographic evaluation of conservation status of African forest  
647 squirrels (Sciuridae) considering land use change and climate change: the importance of point data.  
648 *Biodivers. Conserv.* **16**, 3939–3950 (2007).
- 649 68. Deutsch, C.A. *et al.* Impacts of climate warming on terrestrial ectotherms across latitude. *Proc. Natl.*  
650 *Acad. Sci. U. S. A.* **105**, 6668–72 (2008).
- 651 69. Radchuk, V., Turlure, C. & Schtickzelle, N. Each life stage matters: the importance of assessing the  
652 response to climate change over the complete life cycle in butterflies. *J. Anim. Ecol.* **82**, 275–85  
653 (2013).
- 654 70. Best, A.S., Johst, K., Münkemüller, T. & Travis, M.J. Which species will successfully track climate  
655 change? The influence of intraspecific competition and density dependent dispersal on range shifting  
656 dynamics. *Oikos* **116**, 1531–1539 (2007).
- 657 71. Amstrup, Steven C., Bruce G. Marcot, and David C. Douglas. Forecasting the range wide status of  
658 polar bears at selected times in the 21st century. US Department of the Interior, US Geological  
659 Survey (2007).
- 660 72. Huey, R.B. *et al.* Predicting organismal vulnerability to climate warming: roles of behaviour,  
661 physiology and adaptation. *Philos. Trans. R. Soc. Lond. B. Biol. Sci.* **367**, 1665–79 (2012).
- 662 73. Wilson, R.J., Davies, Z.G. & Thomas, C.D. Modelling the effect of habitat fragmentation on range  
663 expansion in a butterfly. *Proc. Biol. Sci.* **276**, 1421–7 (2009).
- 664 74. Morin, X. & Thuiller, W. Comparing niche- and process-based models to reduce prediction  
665 uncertainty in species range shifts under climate change. *Ecology* **90**, 1301–13 (2009).
- 666 75. Kearney, M. & Porter, W. Mechanistic niche modelling: combining physiological and spatial data to  
667 predict species' ranges. *Ecol. Lett.* **12**, 334–50 (2009).
- 668 76. Monahan, W.B. A mechanistic niche model for measuring species' distributional responses to  
669 seasonal temperature gradients. *PLoS One* **4**, e7921 (2009).
- 670 77. Young, B., Byers, E., Gravuer, K., Hall, K., Hammerson, G., & Redder, A. Guidelines for using the  
671 NatureServe climate change vulnerability index. NatureServe, Arlington, VA (2010).
- 672 78. Gardali, T., Seavy, N.E., DiGaudio, R.T. & Comrack, L.A. A climate change vulnerability  
673 assessment of California's at-risk birds. *PLoS One* **7**, e29507 (2012).
- 674 79. Rowland, E.L., Davison, J.E. & Graumlich, L.J. Approaches to evaluating climate change impacts on  
675 species: a guide to initiating the adaptation planning process. *Environ. Manage.* **47**, 322–37 (2011).
- 676 80. Moyle, P.B., Kiernan, J.D., Crain, P.K. & Quiñones, R.M. Climate change vulnerability of native and  
677 alien freshwater fishes of California: a systematic assessment approach. *PLoS One* **8**, e63883 (2013).
- 678 81. Pacifici, M. *et al.* Generation length for mammals. *Nat. Conserv.* **5**, 89–94 (2013).

- 679 82. Lankford, A.J., Svancara, L.K., Lawler, J.J. & Vierling, K. Comparison of climate change  
680 vulnerability assessments for wildlife. *Wildl. Soc. Bull.* (2014). doi:10.1002/wsb.399
- 681 83. Thomas, C.D. *et al.* A framework for assessing threats and benefits to species responding to climate  
682 change. *Methods Ecol. Evol.* **2**, 125–142 (2011).
- 683 84. Santini, L. *et al.* Ecological correlates of dispersal distance in terrestrial mammals. *Hystrix* **24**, 1–6  
684 (2013).
- 685 85. Maclean, I. M. D. & Wilson, R. J. Recent ecological responses to climate change support predictions  
686 of high extinction risk. *Proc. Natl. Acad. Sci. U. S. A.* **108**, 12337–42 (2011).
- 687 86. IUCN Species Survival Commission. Iucn red list categories and criteria. Version 3.1. (IUCN, Gland,  
688 Switzerland, 2001) (2001).
- 689 87. Akçakaya, H.R., Butchart, S.H.M., Mace, G.M., Stuart, S.N. & Hilton-Taylor, C. Use and misuse of  
690 the IUCN Red List Criteria in projecting climate change impacts on biodiversity. *Glob. Change Biol.*  
691 **12**, 2037–2043 (2006).
- 692 88. Pearson, R.G. *et al.* Life history and spatial traits predict extinction risk due to climate change. *Nat.*  
693 *Clim. Change.* **4**, 217–221 (2014).
- 694 89. Summers, D. M., Bryan, B. a., Crossman, N. D. & Meyer, W. S. Species vulnerability to climate  
695 change: impacts on spatial conservation priorities and species representation. *Glob. Change Biol.* **18**,  
696 2335–2348 (2012).
- 697 90. Rondinini, C. *et al.* Global habitat suitability models of terrestrial mammals. *Philos. Trans. R. Soc.*  
698 *Lond. B. Biol. Sci.* **366**, 2633–41 (2011).
- 699 91. Wintle, B.A.. *et al.* Ecological–economic optimization of biodiversity conservation under climate  
700 change. *Nat. Clim. Change.* **1**, 355–359 (2011).
- 701 92. Thuiller, W., Lavorel, S., Araújo, M.B., Sykes, M.T. & Prentice, I.C. Climate change threats to plant  
702 diversity in Europe. *Proc. Natl. Acad. Sci. U. S. A.* **102**, 8245–50 (2005).
- 703 93. Levinsky, I., Skov, F., Svenning, J.-C. & Rahbek, C. Potential impacts of climate change on the  
704 distributions and diversity patterns of European mammals. *Biodivers. Conserv.* **16**, 3803–3816  
705 (2007).
- 706 94. Poiani, K.A., Goldman, R.L., Hobson, J., Hoekstra, J.M. & Nelson, K.S.. Redesigning biodiversity  
707 conservation projects for climate change: examples from the field. *Biodivers. Conserv.* **20**, 185–201  
708 (2011).
- 709 95. Game, E.T., Kareiva, P. & Possingham, H.P. Six common mistakes in conservation priority setting.  
710 *Conserv. Biol.* **27**, 480–5 (2013).
- 711 96. Bottrill, M.C. *et al.* Is conservation triage just smart decision making? *Trends Ecol. Evol.* **23**, 649–  
712 654 (2008).
- 713 97. Araújo, M.B., Pearson, R.G., Thuiller, W. & Erhard, M. Validation of species–climate impact models  
714 under climate change ´. *Glob. Change Biol.* **11**, 1–10 (2005).
- 715 98. Watling, J.I. *et al.* Validating predictions from climate envelope models. *PLoS One* **8**, e63600 (2013).

- 716 99. Feeley, K.J. & Silman, M.R. Land-use and climate change effects on population size and extinction  
717 risk of Andean plants. *Glob. Change Biol.* **16**, 3215–3222 (2010).
- 718 100. Watson, J.E.M. & Segan, D.B. Accommodating the human response for realistic adaptation planning:  
719 response to Gillson et al. *Trends Ecol. Evol.* **28**, 573–4 (2013).

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727

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729 M.P., P.V., C.R., J.E.M.W. and W.B.F. designed the framework for the review. All authors contributed to the  
730 writing, discussed the results and commented on the manuscript.

731

## 732 **Competing financial interests**

733 The authors declare no competing financial interests.

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735 **Figure headings:**

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737 **Figure 1:** Taxonomic focus of vulnerability assessments in the analysed papers.

738 Birds are the most analysed taxon, followed by mammals and plants, while invertebrates other than  
739 insects have seldom been assessed. Colours represent the spatial scale of the assessments. Regional  
740 scale is defined as describing the range of  $10^4$  to  $10^7$  km<sup>2</sup>, while scales smaller than  $10^4$  km<sup>2</sup> are  
741 referred to as local scales.

742 **Figure 2:** Ecoregional global concentrations of terrestrial and marine climate change vulnerable  
743 species.

744 Studies conducted at regional, continental and global scales were used to derive a global map of  
745 vulnerability, according to an ecoregional classification. The red scale represents terrestrial areas  
746 with high numbers of vulnerable species, identified on the basis of 1) the number of species  
747 assessed and 2) the taxonomic ranks higher than species considered. The blue scale represents areas  
748 that host marine vulnerable species. Dark colours indicate areas of high vulnerability, while light  
749 colours indicate areas of relatively low vulnerability.

750 **Figure 3:** Trends and biases in taxonomic groups assessed and approaches used by continent.

751 Birds and mammals have been the most frequently analysed taxa across all continents between 1995  
752 and 2014, usually with similar proportions (with the exception of Asia). Correlative approaches are  
753 widely used for assessing species vulnerability in Africa, Asia and Europe, while mechanistic  
754 approaches prevail in North America. Trait-based approaches are used mostly in Australia and  
755 North America.

756



**Table 1 | Examples of objectives in climate change vulnerability assessments, on the basis of the scale to be adopted.**

	Temporal scale			Spatial scale			Taxonomic scale		
	Past	Recent past/ present	Present/ Future	Local/site	Regional	Global	Population and ranks < than species	Single species	Multispecies
<b>Examples of objectives: correlative</b>	Reconstructing species' past distribution <sup>101</sup>	Modelling current climatic suitable areas for species <sup>22</sup>	Predicting climate- induced future range shifts under different time intervals <sup>102</sup>	Quantifying the area that will remain climatically suitable for species living in areas important for conservation <sup>60</sup>	Assessing the ability of a network of protected areas to ensure the persistence of species <sup>103</sup>	Identifying the most important climatic variables in determining a species' distribution globally <sup>19</sup>	Quantify the latitudinal/ altitudinal shifts of the various populations of a species <sup>104</sup>	Assessing a species' future threat status <sup>93</sup>	Predicting spatial patterns of species richness <sup>105</sup>
	Identifying past climatic refugia <sup>106</sup>	Quantifying % range gains/losses in the last decades to estimate extinction risk <sup>38</sup>	Projecting future range margin contractions/ expansions by 2080 <sup>92</sup>	Quantifying species' turnover within a protected area <sup>54</sup>	Identifying and designing potential areas to be protected within a region <sup>107</sup>	Identifying hotspots of species highly exposed <sup>19</sup>	Assessing which of the populations of a species will experience the greatest changes in its distribution <sup>104</sup>	Predicting spatial overlap between the current and future range of a species <sup>108</sup>	Modelling possible future community assemblages <sup>109</sup>
<b>Examples of objectives: mechanistic</b>	Representing postglacial expansions from glacial refugia <sup>110</sup>	Quantifying population reductions in recent times due to changes in sea ice extent <sup>41</sup>	Predicting survival under future climate change <sup>111</sup>	Determining climatic factors that affect reproductive success of a reintroduced species <sup>112</sup>	Exploring the range margin dynamics for species of conservation concern within a region <sup>40</sup>	Assessing species thermal tolerances across their range <sup>113</sup>	Assessing the extinction risk of a population at the margins of a species' range <sup>40</sup>	Assessing the impacts of sea level rise on a coastal species <sup>114</sup>	Modelling prey- predator dynamics under future climatic conditions <sup>45</sup>

	Understanding the effects of changes in CO <sub>2</sub> concentration on plants <sup>115</sup>	Determining population viability due to an increase in frequency of extreme climatic events during the last decades <sup>43</sup>	Assessing species' probability of extinction by 2100 <sup>41</sup>	Predicting the probability of extinction of a keystone species within a site <sup>42</sup>	Exploring the extinction risk of a species in part of its range <sup>39</sup>	Predicting changes in fitness due to global warming globally <sup>68</sup>	Determining the extinction risk of a threatened subspecies <sup>34</sup>	Estimating species' abundance in the future under climate change <sup>116</sup>	Predicting future expansions of invasive species <sup>117</sup>
<b>Examples of objectives: TVA</b>	Identifying trends of past extinctions related to life history traits <sup>118</sup>	Identifying taxonomic groups that currently retain high numbers of sensitive and unadaptable species <sup>78</sup>	Identifying sensitive species living in areas that are likely to become highly exposed in the future <sup>119</sup>	Prioritizing conservation actions at the local scale <sup>120</sup>	Making an assessment of species vulnerability within a country <sup>80</sup>	Identifying species with the greatest relative vulnerability to climate change <sup>78</sup>	Identifying potential adaptive characteristics of an isolated subspecies <sup>35</sup>	Identifying the traits that make a species most vulnerable to climate change <sup>120</sup>	Identifying the most vulnerable species to climate change within a taxon <sup>20</sup>
	Predicting the response of species, that share life history traits with past extinct/impacted species, to future climatic changes <sup>121</sup>	Identifying the characteristics of species that played the most important role in determining reductions/ extinctions in recent years <sup>14</sup>	Identifying unadaptable species with the largest predicted range shifts in the coming decades <sup>83</sup>	Understanding which component of vulnerability is prevalent for a species within a site <sup>122</sup>	Understanding how traits relate to changes in occurrence of species within a freshwater basin subject to droughts <sup>11</sup>	Identifying areas with the greatest number of vulnerable species at the global scale <sup>14</sup>	Identifying potentially vulnerable subspecies/ populations/ varieties with relatively unknown distribution <sup>36</sup>	Assessing species' adaptive capacity/ resilience <sup>14</sup>	Selecting different adaptation strategies according to the relative vulnerability of different species <sup>78</sup>

\*References from 101 to 122 are listed in the Supplementary material.

760 **Box 1 | Data availability**

761       Once clear objectives have been established, and the potential approaches identified, another  
762 consideration for selecting the most appropriate method is to consider the types of data available.  
763 The financial resources, time, expertise and input data required for each method are likely to mean  
764 that just one or, at best, a few approaches are feasible. When fine scale data on species occurrence  
765 are available (e.g. point localities), correlative and mechanistic niche models may be applied. To  
766 build these types of models, adequate climate data covering different time periods are also needed.  
767 For example, paleoclimatic reconstructions for Paleocene and Holocene, as well as current and  
768 future projections, are already available under different resolutions and time intervals (e.g.<sup>123,124</sup>).

769       Where relevant life-history data (e.g. data on species' biology, ecology, physiology,  
770 demography) are available; (see ecology and trait databases for birds<sup>125</sup>, mammals<sup>81,126</sup> and  
771 amphibians<sup>127</sup>), trait-based or mechanistic approaches could facilitate, for example, the  
772 identification of resilient and/or adaptable species, thus aiding in prioritization<sup>11</sup>. Moreover, these  
773 kinds of data are necessary to develop mechanistic niche models to refine species' distribution based  
774 on the mechanisms that species themselves develop to cope with global warming<sup>13</sup>. Often this type  
775 of empirical data will be lacking. Rather than abandon modelling and informing conservation  
776 decisions in these cases, structured expert elicitation approaches offer an interim way of estimating  
777 key species demographic and life-history parameters<sup>128,129</sup>.

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